

Polar Ozone Loss in a Changing Climate

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SUMMARY

The goal of this work is to robustly simulate polar ozone (O_3) variability during recent years by optimizing a version of the passive ozone subtraction technique. The eventual goal is to predict future polar O₃ loss in a changed climate, and to explore how the atmosphere responds to polar ozone recovery. The passive O₃ subtraction technique subtracts simulated, inert (or partially inert) O₃ from observed or predicted ozone. The work here uses the Specified Dynamics Whole Atmosphere Community Climate Model (SD-WACCM) to simulate a "pseudo"-passive O₃ tracer which only non-halogen chemistry is allowed to perturb. Observations are from the Aura Microwave Limb Sounder (MLS). O₃ loss calculations during Arctic winter 2004/05 are in good agreement with previous work, providing an initial verification that SD-WACCM is appropriate for these types of studies. Diagnostic comparisons to observations of ozone-related species point to minor deficiencies in SD-WACCM simulations of descent and/or mixing, as well as halogen-induced O_3 depletion.

METHOD

- Three model simulations:
- full-ozone chemistry
- gas-phase-ozone chemistry only (pseudo-passive tracer)
- no ozone chemistry (passive tracer, for reference)
- Inferred Loss: O_3 loss quantified by model & measurement $IL = (EOS MLS O_3) (SD-WACCM pseudo-passive <math>O_3$)
- Modeled Loss: O_3 loss quantified by model only $ML = (SD-WACCM O_3) (SD-WACCM pseudo-passive <math>O_3)$

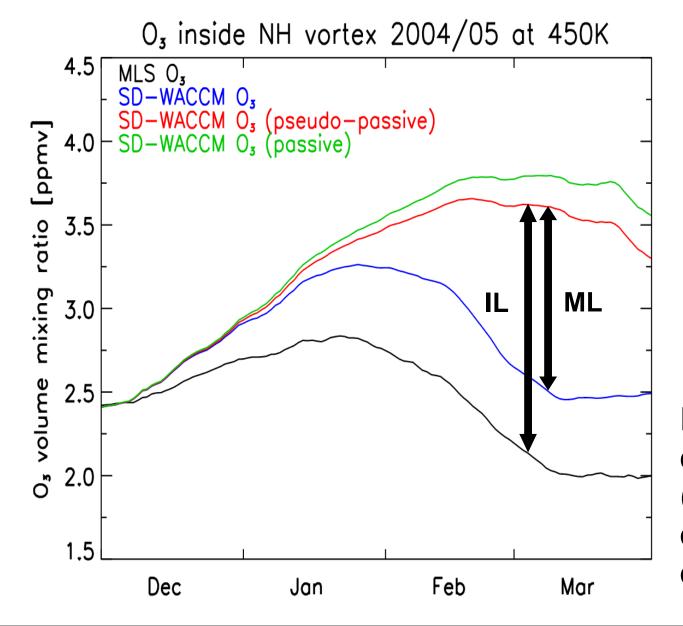


Figure 1: Evolution of observed ozone (black), modeled ozone (blue), modeled pseudo-passive ozone (red), and modeled passive ozone (green).

DATA

- EOS MLS on Aura (since August 2004), version 2.2
- SD-WACCM (nudged daily with GEOS* U, V, T), version 3548
 *Goddard Earth Observing System, reanalysis, version 5

INITIALIZATION

Global O_3 , nitrous oxide (N_2O) , nitric acid (HNO_3) , hydrogen chloride (HCI), and water vapor (H_2O) initialized with MLS data

- On 1 Dec: before first O₃ loss occurs
- MLS data interpolated to SD-WACCM grid
- Interpolation done on SD-WACCM pressure levels
- Delaunay-Triangulation
- Equal-area smoothing
- Cannot treat diurnal variations (e.g. chlorine monoxide (CIO))

REFERENCES

Jin, J. J., et al. (2006), *Geophys. Res. Lett.*, 33, L15801, doi:10.1029/2006GL026752. Manney, G. L., et al. (2006), *Geophys. Res. Lett.*, 33, L04802, doi:10.1029/2005GL024494. Rex, M., et al. (2006), *Geophys. Res. Lett.*, 33, L23808, doi:10.1029/2006GL026731. Rösevall, J. D., et al. (2008), *J. Geophys. Res.*, 113, D13301, doi:10.1029/2007JD009560. Santee, M. L., et al. (2008), *J. Geophys. Res.*, 113, D12307, doi:10.1029/2007JD009057. Singleton, C. S., et al. (2007), *J. Geophys. Res.*, 112, D07304, doi:10.1029/2006JD007463.

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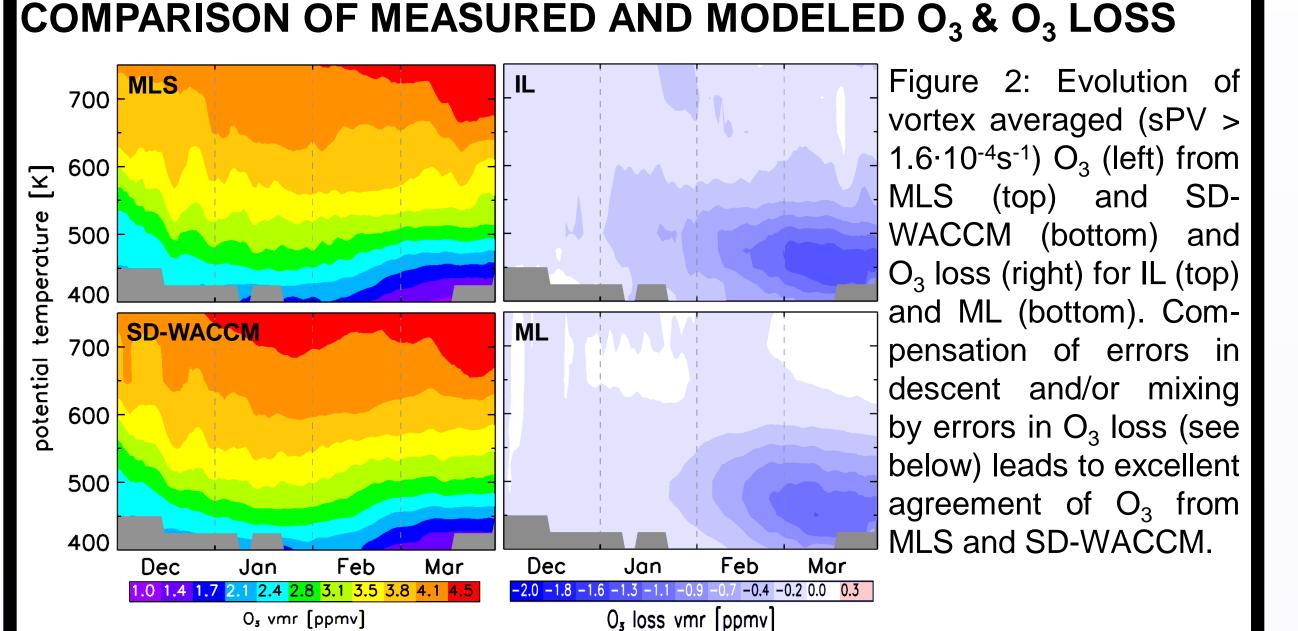


Figure 3: Spatial distribution of inferred O_3 loss at 450K in 10day intervals throughout the season (same color bar as Fig. 2) with a green 1.6·10⁻⁴s⁻¹ sPV contour. Largest O_3 loss of 2ppmv occurs at the end of the season.

Table 1: Comparison of shown O_3 loss results (last column) with previous research. Potential Temperature | Manney et al. [2006] | Jin et al. [2006] | Rex et al. [2006] | Rösevall et al. [2008] | Singleton et al. [2007] | MLS/WACCM $1.6 \pm 0.3 \text{ ppmv}$ 0.7 ppmv 0.8 ppmv 1.4 ppmv 1.3 ppmv 1.2 - 1.5 ppmv 2.2 ppmv 2.0 ppmv $1.7 \pm 0.4 \text{ ppmv}$ 1.3 ppmv 1.2 - 1.5 ppmv 2.1 ppmv $1.1 \pm 0.4 \text{ ppmv}$ 0.8 ppmv 1.8 ppmv 1.2 ppmv $0.6 \pm 0.3 \text{ ppmv}$ 0.5 ppmv 1.3 ppmv 0.6 ppmv 0.6 ppmv 0.4 ppmv

COMPARISON OF MEASURED AND MODELED N₂O Figure 4 (left): Evolution of MLS (top) and SD-WACCM (bottom) vortex averaged N₂O (sPV > 1.6·10⁻⁴s⁻¹). Differences indicate errors in SD-WACCM simulation of descent and/or mixing. However, no clear distinction between differences in descent and mixing across the vortex edge can be Figure 5 (below): Spatial distribution of N₂O (1st row MLS, 2nd row SD-WACCM) at 490K for one day 600 each month throughout the season with a 1.6·10⁻⁴s⁻¹ sPV line contour. The final warming happened ~3/10 [Manney et al., 2006]. Differences between SD-WACCM and MLS N₂O mixing ratios suggest less descent inside polar vortex for SD-WACCM compared to MLS. 03/15/2004 01/29/2004 12/23/2004 02/27/2004

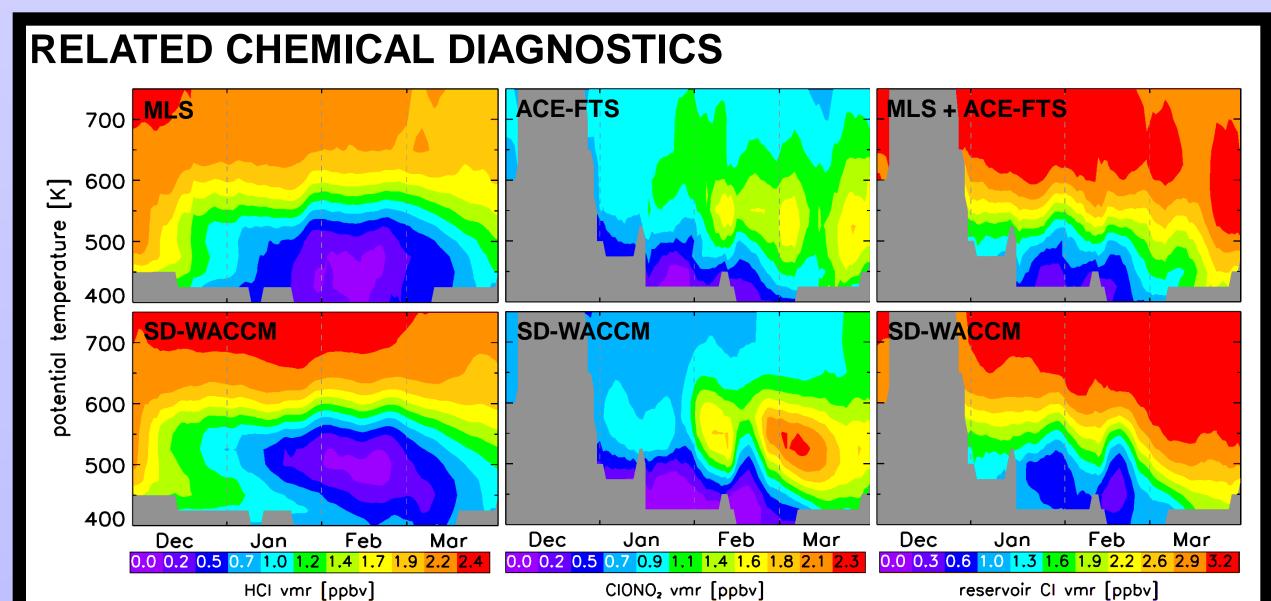


Figure 6: Evolution of observed (top row) HCl (left), ClONO₂ (middle), their sum (right) and SD-WACCM respectively (bottom row) inside the polar vortex (sPV > $1.6 \cdot 10^{-4} \text{s}^{-1}$). Cl in reservoir species from SD-WACCM compares well with observations, suggesting the correct partitioning of chlorine between reactive forms and reservoirs.

RELATED MICROPHYSICAL DIAGNOSTICS

Figure 7 (left): Evolution of polar vortex averaged (sPV > $1.6\cdot10^{-4}$ s⁻¹) gas-phase HNO₃ from SD-WACCM (bottom) compares well with MLS (top). Slight underestimates in the model are found later in the season below ~550K.

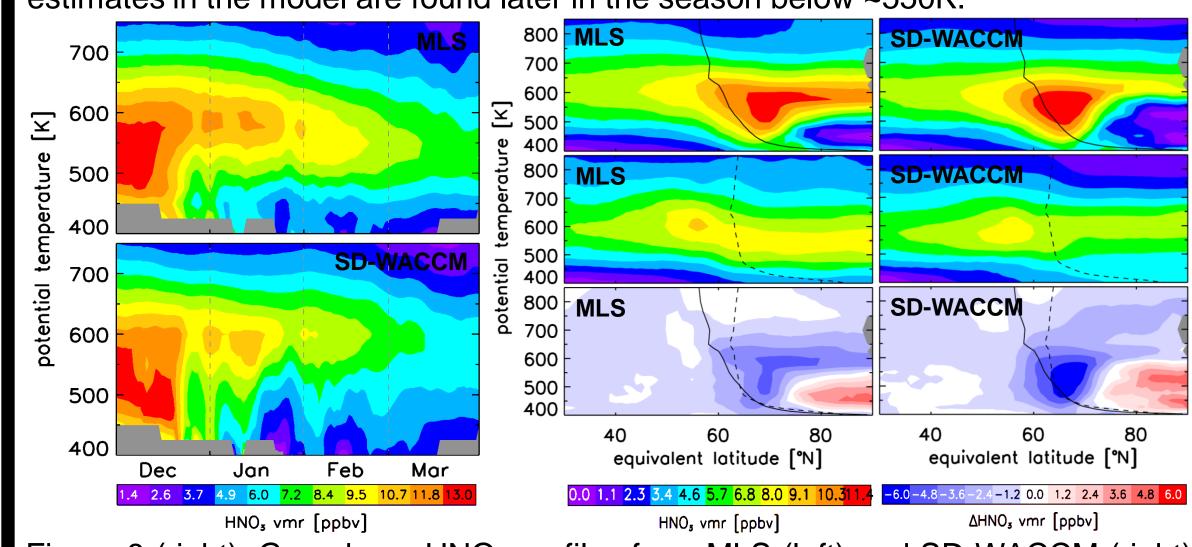
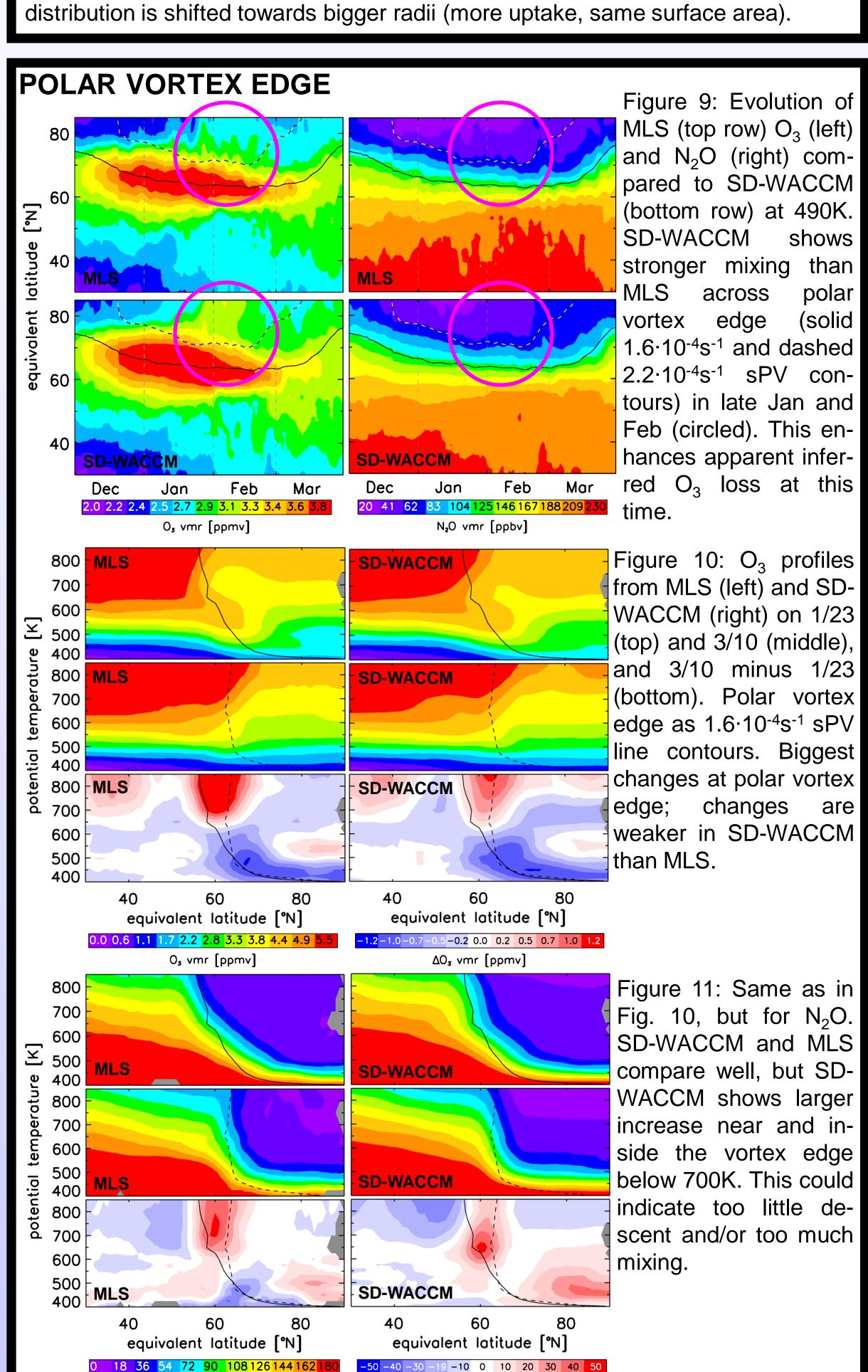


Figure 8 (right): Gas-phase HNO₃ profiles from MLS (left) and SD-WACCM (right) on 1/23 (top) and 3/10 (middle), and 3/10 minus 1/23 (bottom).

Too much uptake of gas-phase HNO_3 in SD-WACCM is consistent with too little $CIONO_2$ (see Fig. 6). Reasonable O_3 loss then suggests that PSC particle size distribution is shifted towards bigger radii (more uptake, same surface area).



CONCLUSIONS

N₂O vmr [ppbv]

- SD-WACCM is valid for inferring O₃ loss from observations
- More accurate simulation of O₃ loss in WACCM requires further investigation of chlorine partitioning and PSC particle sizes

ΔN₂O vmr [ppbv]

- Equivalent analysis for Antarctic winter needed to better investigate mixing and descent
- Future plans include O_3 loss calculations for all Arctic and Antarctic winters since 2004